AMENDMENT

Kindly amend the application, without prejudice, as follows:

IN THE CLAIMS:

Kindly amend claims 1-16, without prejudice, to read as follows:

- 1. (Currently Amended) A Mmethod of quantifying viral or bacterial particles having a cholesterol-containing envelope, comprising the steps of: staining wherein the particles are stained with a fluorogenic polyene macrolide and quantitatively determining the fluorescence signals of the individual particles by counting the fluorescence signals under a fluorescence microscope are then quantitatively determined to provide a particle number.
- 2. (Currently Amended) The mMethod according to claim 1, wherein the method is applied to retroviruses, orthomyxoviruses, paramyxoviruses, togaviruses, bunyaviruses, rhabdoviruses, filoviruses, arenoviruses, coronaviruses, herpesviruses, flaviviruses, hepadnaviruses, poxviruses or irdoviruses.
- 3 (Currently Amended) The mMethod according to claim 1, wherein the method is applied to HIV, measles virus, influenza virus, murine leukaemia virus or mycoplasmas.
 - 4. (Canceled).
- 5. (Currently Amended) The mMethod according to claim 1, wherein the polyene macrolide is filipin is used as the polyene macrolide.
- 6. (Currently Amended) The mMethod according to claim 5, wherein the filipin fluorescence is excited at a wavelength of 387 ± 14 nm and the counting is carried out at the emission wavelength of 450 ± 29 nm.
- 7. (Currently Amended) The mMethod according to claim 1, wherein for quantitative determination, the number and/or concentration of fluorescent particles is compared

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to the <u>a</u> known number and/or concentration of specified fluorescent particles <u>as a reference</u> standard.

- 8. (Currently Amended) The mMethod according to claim 7, wherein for eomparison, the fluorescent particles as the reference standard are specified that are from 0.5 times to twice as large as, especially, about the same size as, the particles being quantified.
- 9. (Currently Amended) The mMethod according to claim 7, wherein-for eomparison, the inert fluorescent particles as the reference standard are specified are inert fluorescent particles.
- 10. (Currently Amended) A kKit of parts for quantifying viral or bacterial particles having a cholesterol-containing envelope, which comprises
 - a fluorogenic polyene macrolide and

 (as optional constituent) fluorescent particles as a reference standard.
- 11. (Currently Amended) The kKit of parts according to claim 10, wherein the reference standard is present in an aqueous medium.
- 12. (Currently Amended) The kKit of parts according to claim 10, wherein the fluorescent particles of the reference standard are inert particles.
- 13. (Currently Amended) The kKit of parts according to claim 10, wherein the fluorescent particles of the reference standard are from 0.5 times to twice as large as, and especially about the same size as, the particles being quantified.
- 14. (Currently Amended) The kKit of parts according to claim 10, wherein the polyene macrolide is having filipin as the polyene macrolide.
- 15. (Currently Amended) A method of using Use of a fluorogenic polyene macrolide especially filipin, for quantifying viral particles having a cholesterol-containing envelope

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comprising the steps of: staining the viral particle with the fluorogenic polyene macrolide and quantifying fluorescence signals of the viral particles under a fluorescence microscope.

- 16 (New) The method according to claim 8, wherein the fluorescent particles reference standard is about the same size as the particles being quantified.
- 17. (New) The kit of parts according to claim 13, wherein the fluorescent particles of the reference standard are about the same size as the particles being quantified.
- 18. (New) The method according to claim 15, wherein the polyene macrolide is filipin.

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